Refine Search

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Terms	Documents			
(cepa or fistulosum) and L10	0			

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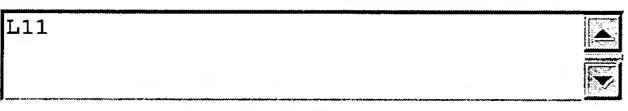
EPO Abstracts Database

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<u>L3</u>	(embryogen\$ and call\$) and L2	21	<u>L3</u>					
<u>L2</u>	(cepa or fistulosum) and L1	47	<u>L2</u>					
L1	alli\$ and agrobacter\$	452	L1					

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JAN 17 IPC 8 in the WPI family of databases including WPIFV

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Transgenic rose lines harboring an antimicrobial protein gene, Ace-AMP1,

 ${ t TI}$

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demonstrate enhanced resistance to powdery mildew (Sphaerotheca pannosa)
     Li, Xianggian; Gasic, Ksenjia; Cammue, Bruno; Broekaert, Willem; Korban,
AU
     Schuyler S.
     Department of Natural Resources and Environmental Sciences, University of
CS
     Illinois, Urbana, IL, 618001, USA
     Planta (2003), 218(2), 226-232
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     Springer-Verlag
PB
     Journal
DT
     English
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AN
     2002:977962 CAPLUS
DN
     138:36240
     Improved efficiency of regeneration of transgenic plants using
TI
     meristematic or nodal tissue transformed with Agrobacterium
     Goldman, Stephen L.; Rudrabhatla, Sairam V.
IN
     University of Toledo, USA
PA
     PCT Int. Appl., 84 pp.
SO
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AN
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DN
     Process for inducing direct somatic embryogenesis and secondary
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     embryogenesis in monocotyledonous plant cells, and rapidly
     regenerating fertile plants
     Eudes, Francois Andre Germain; Laroche, Andre J.; Acharya, Surya Narayan
IN
     Her Majesty the Queen in Right of Canada as Represented by the Minister of
PA
     Agriculture and Agri-Food, Can.
     PCT Int. Appl., 70 pp.
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PATENT NO.

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APPLICATION NO.

DATE

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     137:349439
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     Process for inducing direct somatic embryogenesis and secondary
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     embryogenesis in monocotyledonous plant cells, and rapidly
     regenerating fertile plants
     Eudes, Francois Andre Germain; Laroche, Andre J.; Acharya, Surya Narayan
IN
PA
     U.S. Pat. Appl. Publ., 31 pp., Cont.-in-part of U.S. Ser. No. 641,243.
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L4
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     2004:962821 CAPLUS
DN
     142:194462
TI
     Process for inducing direct somatic embryogenesis and secondary
     embryogenesis in monocotyledonous plant cells, and rapidly
     regenerating fertile plants
     Eudes, Francois Andre Germain; Acharya, Surya Narayan; Laroche, Andre J.
IN
     Her Majesty the Queen In Right of Canada as Represented by the Minister,
PΑ
     Can.
    Can. Pat. Appl., 60 pp.
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    ANSWER 8 OF 11 CABA COPYRIGHT 2006 CABI on STN
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     2002:204189 CABA
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     20023153005
    Embryogenic callus induction from leaf explants of the
TI
    Liliaceous ornamental plant, Agapanthus praecox ssp. orientalis (Leighton)
    Leighton histological study and response to selective agents
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Faculty of Agriculture, Niigata University, 2-8050 Ikarashi, Niigata

Suzuki, S.; Oota, M.; Nakano, M.

950-2181, Japan. mnakano@agr.niigata-u.ac.jp

AU

CS

Scientia Horticulturae, (2002) Vol. 95, No. 1/2, pp. 123-132. 24 ref. SO Publisher: Elsevier Science B.V. Amsterdam ISSN: 0304-4238 DOI: 10.1016/S0304-4238(02)00033-X Netherlands Antilles CY Journal DTEnglish LA EDEntered STN: 20021206 Last Updated on STN: 20021206 ANSWER 9 OF 11 CABA COPYRIGHT 2006 CABI on STN L42003:30105 CABA AN 20023198899 DN Agrobacterium-mediated transformation in Liliaceous ornamental \mathtt{TI} plants Suzuki, S.; Nakano, M. AU Faculty of Agriculture, Niigata University, 2-8050 Ikarashi, Niigata CS 950-2181, Japan. mnakano@agr.niigata-u.ac.jp JARQ, Japan Agricultural Research Quarterly, (2002) Vol. 36, No. 3, pp. SO 119-127. 26 ref. Publisher: Japan International Research Center for Agricultural Sciences. Tsukuba ISSN: 0021-3551 CY Japan Journal \mathtt{DT} English LA Entered STN: 20030214 EDLast Updated on STN: 20030214 ANSWER 10 OF 11 CABA COPYRIGHT 2006 CABI on STN L4AN 2001:104509 CABA 20013082803 DNProduction of transgenic plants of the Liliaceous ornamental plant TIAgapanthus praecox ssp. orientalis (Leighton) Leighton via Agrobacterium-mediated transformation of embryogenic calli Suzuki, S.; Supaibulwatana, K.; Mii, M.; Nakano, M.; Kanyaratt AU Supaibulwatana Faculty of Agriculture, Niigata University, 2-8050 Ikarashi, Niigata CS 950-2181, Japan. Plant Science, (2001) Vol. 161, No. 1, pp. 89-97. 29 ref. SO Publisher: Elsevier Science Ltd. Oxford ISSN: 0168-9452 United Kingdom CYJournal DTEnglish LAED Entered STN: 20011004 Last Updated on STN: 20011004 L4ANSWER 11 OF 11 CAPLUS COPYRIGHT 2006 ACS on STN AN2000:790242 CAPLUS DN 133:330528 Transformation of Allium sp. with agrobacterium using TIembryogenic callus cultures Reynolds, John IN Seminis Vegetable Seeds, Inc., USA PA PCT Int. Appl., 22 pp. SO CODEN: PIXXD2 \mathtt{DT} Patent English LA FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE ----

20001109 WO 2000-US12463 20000505

WO 2000065903 A1

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ANSWER 1 OF 11 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN L4

2005:518465 BIOSIS ACCESSION NUMBER: DOCUMENT NUMBER: PREV200510305997

Screening wheat genotypes for high callus TITLE:

induction and regeneration capability from immature embryo

cultures.

Haliloglu, Kamil [Reprint Author]; Baenziger, P. Stephen AUTHOR (S): Ataturk Univ, Fac Agr, Dept Field Crops, TR-25240 Erzurum, CORPORATE SOURCE:

Turkey

kamilh@atauni.edu.tr

Journal of Plant Biochemistry and Biotechnology, (JUL 2005) SOURCE:

Vol. 14, No. 2, pp. 155-160.

ALL CITATIONS AVAILABLE IN THE RE FORMAT

ISSN: 0971-7811.

DOCUMENT TYPE:

Article English LANGUAGE:

ENTRY DATE: Entered STN: 23 Nov 2005

Last Updated on STN: 23 Nov 2005

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BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN L4ANSWER 1 OF 11

2005:518465 BIOSIS ACCESSION NUMBER: PREV200510305997 DOCUMENT NUMBER:

Screening wheat genotypes for high callus TITLE:

induction and regeneration capability from immature embryo

cultures.

Haliloglu, Kamil [Reprint Author]; Baenziger, P. Stephen AUTHOR(S):

CORPORATE SOURCE: Ataturk Univ, Fac Agr, Dept Field Crops, TR-25240 Erzurum,

Turkey

kamilh@atauni.edu.tr

SOURCE: Journal of Plant Biochemistry and Biotechnology, (JUL 2005)

Vol. 14, No. 2, pp. 155-160.

ISSN: 0971-7811.

DOCUMENT TYPE:

Article English

LANGUAGE: ENTRY DATE:

Entered STN: 23 Nov 2005

Last Updated on STN: 23 Nov 2005

Selecting the explant genotypes is crucial step in in vitro culture and AB Agrobacterium-mediated transformation system due to its host range Immature embryos of five winter and three spring wheat (Triticum aestivum L) cultivars were evaluated for tissue culture response in three callus initiation media. MS medium containing 2,4-D (2

mg ml(-1)) plus B5 vitamins (MSB5), MS medium containing 2,4-D (1 mg ml(-1)) with no vitamins (MS1GC) or MS medium containing picloram (2.2 mg ml(-1)) and 2,4-D (0.5 mg ml(-1)) plus MS vitamins (CM4C) were used for callus initiation. Percentage of callus induction varied widely with the genotype and initiation medium used, with values ranging from 5.7% to 100%. Embryogenic capacity of genotypes was evaluated by number of somatic embryos formed from cultured immature embryos. Bob White (spring) and NE92458 (winter) were equal and most embryogenic; Pronghorn and 2137 (both winter) were the poorest. CM4C medium was found to be the best medium for initiating embryogenic callus among three culture media tested. A standard, regeneration procedure was used. The genotypes with the highest regeneration efficiencies were Bob White, Fielder and NE92458, (1.8, 1.4 and 1.6 plants/explant, respectively).

L4 ANSWER 2 OF 11 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN

ACCESSION NUMBER: 2005:43507 BIOSIS DOCUMENT NUMBER: PREV200500043533

TITLE: Genetic transformation of garlic (Allium sativum

L.) by particle bombardment.

AUTHOR(S): Robledo-Paz, Alejandrina [Reprint Author]; Cabrera-Ponce,

Jose Luis; Villalobos-Arambula, Victor Manuel; Herrera-Estrella, Luis; Jofre-Garfias, Alba Estela

CORPORATE SOURCE: Ctr Invest and Estudios AvanzadosDept Ingn Genet Plantas,

IPN, Km 9-6 Libramiento Norte Carretera Irapuato Leon,,

Irapuato, Gto, 36500, Mexico

arobledo@colpos.mx

SOURCE: Hortscience, (October 2004) Vol. 39, No. 6, pp. 1208-1211.

print.

ISSN: 0018-5345 (ISSN print).

DOCUMENT TYPE: Article

LANGUAGE: English

ENTRY DATE: Entered STN: 26 Jan 2005

Last Updated on STN: 26 Jan 2005

AB Microprojectile bombardment was used to introduce DNA into

embryogenic callus of garlic (Allium sativum

L.) and produce stably transformed garlic plants. Embryogenic calluses, derived from garlic cultivar 'GT96-1', were bombarded with plasmid DNA containing genes coding for hygromycin phosphotransferase and beta-glucuronidase. Putatively transformed calluses were identified in the bombarded tissue after 4 months of selection on 20 mg.L-1 hygromycin B. The transgenic nature of the selected material was demonstrated by GUS histochemical assay and Southern blot hybridization analysis, and twenty transgenic plants were regenerated.

L4 ANSWER 3 OF 11 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 1

ACCESSION NUMBER: 2003:933684 CAPLUS

DOCUMENT NUMBER: 140:175930

TITLE: Transgenic rose lines harboring an antimicrobial

protein gene, Ace-AMP1, demonstrate enhanced

resistance to powdery mildew (Sphaerotheca pannosa)

AUTHOR(S): Li, Xiangqian; Gasic, Ksenjia; Cammue, Bruno;

Broekaert, Willem; Korban, Schuyler S.

CORPORATE SOURCE: Department of Natural Resources and Environmental

Sciences, University of Illinois, Urbana, IL, 618001,

USA

SOURCE: Planta (2003), 218(2), 226-232

CODEN: PLANAB; ISSN: 0032-0935

PUBLISHER: Springer-Verlag

DOCUMENT TYPE: Journal LANGUAGE: English

AB An antimicrobial protein gene, Ace-AMP1, was introduced into Rosa hybrida cv. Carefree Beauty via Agrobacterium-mediated transformation.

A total of 500 putative transgenic plants were obtained from 100 primary

embryogenic calli co-cultivated with A. tumefaciens following selection on a regeneration medium containing 100 mg/l kanamycin. Polymerase chain reaction anal. of these putative transgenic lines, using primers for both Ace-AMP1 and neomycin phosphotransferase (npt II) genes, showed that 62% of these plants were pos. for both transgenes. These lines were further confirmed for stable integration of Ace-AMP1 and npt II genes by Southern blotting. Transcription of the Ace-AMP1 transgene in various transgenic rose lines was determined using Northern blotting. Transgenic rose lines inoculated with conidial spores of Sphaerotheca pannosa (Wallr.: Fr.) Lev. var. rosae showed enhanced resistance to powdery mildew using both a detached-leaf assay and an in vivo greenhouse whole-plant assay.

REFERENCE COUNT: 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 4 OF 11 CAPLUS COPYRIGHT 2006 ACS on STN L4

ACCESSION NUMBER:

2002:977962 CAPLUS

DOCUMENT NUMBER:

138:36240

TITLE:

Improved efficiency of regeneration of transgenic plants using meristematic or nodal tissue transformed

with Agrobacterium

INVENTOR(S):

Goldman, Stephen L.; Rudrabhatla, Sairam V.

PATENT ASSIGNEE(S):

University of Toledo, USA

SOURCE:

PCT Int. Appl., 84 pp. CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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			GR,	ΙE,	IT,	LU,	MC,	NL,	PT,	SE,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GA,
			GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG							
	CA	2451	863			AA		2002	1227	(CA 2	002-2	2451	863		2	0020	514
	EP	1455	568			A2		2004	0915]	EP 2	002-	7421	06		2	0020	514
		R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,
			ΙE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	AL,	TR						
	US	2004	2371	3 3		A1		2004	1125	1	US 2	003-4	48086	65		20	0031	212
PRIORITY APPLN. INFO.: US 2001-298542P P 20010615							515											
										1	US 2	002-3	35656	63P	I	2 (0020	211
										1	WO 2	002-1	JS18	966	V	1 2	0020	514
2 10	34-4		_ &	- E E L								7 7				•	1 _ 4	

AB Methods of efficiently transforming monocotyledonous and dicotyledonous plant tissue and regenerating plants with a very high yield of transgenic plants are described. The method uses Agrobacterium to transform root or apical meristem that is then cultured under conditions that generate somatic embryogenesis. The time required for the production of transgenic plants is significantly decreased, while the number of transgenic plants is significantly increased. These increases are not dependent upon the use of super-virulent Agrobacterium strains. The invention also relates to an improved technique for in vitro regeneration of mono- and di-cotyledonous plants in a suitable medium containing a novel growth regulator regime that promotes cell elongation in

the production of numerous somatic embryos that are regenerable into fertile plants. Optimization expts. for the transformation of grasses and legumes using a β-glucuronidase reporter gene are described. Efficient genotype-independent regeneration of transgenic corn is demonstrated.

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ANSWER 5 OF 11 CAPLUS COPYRIGHT 2006 ACS on STN
L4
ACCESSION NUMBER:
                        2002:142892 CAPLUS
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DOCUMENT NUMBER: 136:180165

Process for inducing direct somatic TITLE:

embryogenesis and secondary

embryogenesis in monocotyledonous plant cells,

and rapidly regenerating fertile plants

Eudes, Francois Andre Germain; Laroche, Andre J.; INVENTOR(S):

Acharya, Surya Narayan

Her Majesty the Queen in Right of Canada as PATENT ASSIGNEE(S):

Represented by the Minister of Agriculture and

Agri-Food, Can.

PCT Int. Appl., 70 pp. SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE:

1

Patent English

LANGUAGE: FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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APPLICATION NO. DATE
     PATENT NO.
                      KIND
                               DATE
                                           WO 2001-CA1165
     WO 2002014520
                         A2
                                20020221
                                                                  20010817
     WO 2002014520
                         A3
                                20030213
            AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
            LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL,
             PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG,
            US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
        RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
            DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
            BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                               20020225
                         A5
    AU 2001091535
                                           AU 2001-91535
                                                                   20010817
                                           US 2000-641243 A 20000817
WO 2001-CA1165 W 20010817
PRIORITY APPLN. INFO.:
```

A process for inducing direct somatic embryogenesis and ABsecondary embryogenesis in monocotyledonous plant cells and rapidly regenerating fertile monocotyledonous plants is provided. provided is a process for inducing direct somatic embryogenesis in monocotyledonous plant cells and rapidly regenerating fertile monocotyledonous plants without secondary embryogenesis. Also provided is a process for inducing direct somatic embryogenesis and organogenesis in monocotyledonous plant cells and rapidly regenerating fertile monocotyledonous plants. Also provided is a process for inducing somatic embryogenesis in monocotyledonous callus cells, suspension cells, or microspore-derived embryos, and rapidly regenerating fertile monocotyledonous plants. In contrast to prior art tissue culture methods involving indirect somatic embryogenesis, direct somatic embryogenesis avoids a callus step, and its attendant problems, such as increased somaclonal variation. Tissue culture steps of the invention progress on the basis of the developmental stage of the cultured cells, rather than in accordance with a pre-determined time line, thereby providing green, fertile plants more rapidly than do previous culture methods. In the first step, embryogenic monocotyledonous plant cells are cultured under conditions conductive to direct formation of primary embryos without an intervening callus stage; the cells are not cultured for a pre-determined period of time, but rather until a desired developmental stage is detected. In a second step, one or more of the globular-stage primary embryos from the first step are

cultured under conditions conducive to induction of secondary embryo formation, at least until secondary embryogenesis is detected. In the third step, the one or more secondary embryos from the second step are cultured under conditions conducive to regeneration of plantlets from the secondary embryos. The direct somatic embryogenesis method for monocots provides for the ready introduction of foreign genes into the plant.

L4 ANSWER 6 OF 11 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2002:850253 CAPLUS

DOCUMENT NUMBER: 137:349439

TITLE: Process for inducing direct somatic

embryogenesis and secondary

embryogenesis in monocotyledonous plant cells,

APPLICATION NO.

DATE

and rapidly regenerating fertile plants

INVENTOR(S): Eudes, François Andre Germain; Laroche, Andre J.;

Acharya, Surya Narayan

PATENT ASSIGNEE(S): Can.

SOURCE: U.S. Pat. Appl. Publ., 31 pp., Cont.-in-part of U.S.

Ser. No. 641,243.

DATE

CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

English

KIND

FAMILY ACC. NUM. COUNT: 3

PATENT INFORMATION:

PATENT NO.

	TAILNI NO.	KIND	DAIL	AITEICHTION NO.	DHIB
PR	US 2002164798 IORITY APPLN. INFO.:	A1	20021107	US 2001-929831 US 2000-641243	20010814 A2 20000817
PRAB	A process for indusecondary embryogerapidly regenerations that step, embryo reactions are detected inducing direct somatic embryogenesis. In monocotyledonous plantlets from the first step are developmental stage, at least under conditions conducing direct somatic embryogenesis. In monocotyledonous plantlets from the inducing direct somation of stage, at least undevelopmental stage monocotyledonous plantlets from the inducing direct somation of stage are cultured from the inducing direct something direct somethi	enesis in a first a fi	rect somatic monocotyle ile monocotyle or conducive lus stage, e globular in a second ed under conditions can also problem in monocotyle or conditions can also problem in monocotyle embryos ile are cultiple embryos ile and rapiduction of cor more of expension at ic embryos ile embryos	US 2000-641243 c embryogenesis and edonous plant cells are yledonous plants is proposed formation of at least until at least developmental stage and step, one or more primalitions conducive to condary embryogenesis ore secondary embryos onducive to regeneration of a process for cotyledonous plant celly yledonous plants, with tryogenic tured under conditions without an intervening rimary embryo reaches ep, one or more primary enditions conducive to Also provided is a prosess ep, one or more primary enditions conducive to also provided is a prosess ep, one or more primary enditions conducive to also provided is a prosess end organogenesis in idly regenerating fertitage embryos obtained sis are cultured under organogenesis, or untit the new shoots are the ration of plantlets. Ogenesis in monocotyle icrospore-derived embry yledonous plants. In	A2 20000817 In a sovided. In a sprimary embryos st one induction of is from the second on of plantlets or inducing als and sout secondary Is conducive to callus the globular sy embryos from regeneration of coess for a sile by this same all adventitious and alous routed Also provided is adonous roos, and
	cerrs or mrcrospor	e-derive	ed elimitans	are cultured in or on	a culture mealum

comprising auxin, cytokinin, and polyamine in amts. effective to cause induction of embryo formation, the cytokinin being present in greater proportion than the auxin, at least until at least one embryo reaches the globular developmental stage. In a second step, one or more globular-stage embryos from the first step are cultured under conditions conducive to regeneration of plantlets from the globular-stage embryos. Fertile monocotyledonous plants produced according to the processes of the invention are also provided.

L4 ANSWER 7 OF 11 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2004:962821 CAPLUS

DOCUMENT NUMBER: 142:194462

TITLE: Process for inducing direct somatic

embryogenesis and secondary

embryogenesis in monocotyledonous plant cells,

and rapidly regenerating fertile plants

INVENTOR(S): Eudes, Francois Andre Germain; Acharya, Surya Narayan;

Laroche, Andre J.

PATENT ASSIGNEE(S): Her Majesty the Queen In Right of Canada as

Represented by the Minister, Can.

SOURCE: Can. Pat. Appl., 60 pp.

CODEN: CPXXEB

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 3

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE		
						
CA 2316106	AA	20020217	CA 2000-2316106		20000817	
CA 2355340	AA	20020217	CA 2001-2355340		20010815	
PRIORITY APPLN. INFO.:			CA 2000-2316106	Α	20000817	
			US 2000-641243	A	20000817	

A process for inducing direct somatic embryogenesis and AB secondary embryogenesis in monocotyledonous plant cells and rapidly regenerating fertile monocotyledonous plants is provided. In a first step, embryogenic monocotyledonous plant cells are cultured under conditions conducive to direct formation of primary embryos without an intervening callus stage, at least until at least one primary embryo reaches the globular developmental stage and no longer than the coleoptilar stage. In a second step, one or more primary embryos from the first step are cultured under conditions conducive to induction of secondary embryo formation, until secondary embryogenesis is detected. In a third step, one or more secondary embryos from the second step are cultured under conditions conducive to regeneration of plantlets from the secondary embryos. Also provided is a process for inducing direct somatic embryogenesis in monocotyledonous plant cells and rapidly regenerating fertile monocotyledonous plants, without secondary embryogenesis. In a first step, embryogenic monocotyledonous plant cells are cultured under conditions conducive to direct formation of primary embryos without an intervening callus stage, at least until at least one primary embryo reaches the globular developmental stage. In a second step, one or more primary embryos from the first step are cultured under conditions conducive to regeneration of plantlets from the primary embryos. Also provided is a process for inducing somatic embryogenesis in monocotyledonous callus cells, suspension cells, or microspore-derived embryos, and rapidly regenerating fertile monocotyledonous plants. In a first step embryogenic monocotyledonous callus cells, suspension cells or microspore- derived embryos are cultured in or on a culture medium comprising auxin, cytokinin, and polyamine in amts. effective to cause induction of embryo formation, the cytokinin being present in greater proportion than the auxin, at least until at least one embryo reaches the globular developmental stage. In a second step, one or more

globular-stage embryos from the first step are cultured under conditions conducive to regeneration of plantlets from the globular- stage embryos. Fertile monocotyledonous plants produced according to the processes of the invention are also provided.

L4 ANSWER 8 OF 11 CABA COPYRIGHT 2006 CABI on STN

ACCESSION NUMBER: 2002:204189 CABA

DOCUMENT NUMBER: 20023153005

TITLE: Embryogenic callus induction

from leaf explants of the Liliaceous ornamental plant, Agapanthus praecox ssp. orientalis (Leighton)

Leighton histological study and response to

selective agents

AUTHOR: Suzuki, S.; Oota, M.; Nakano, M.

CORPORATE SOURCE: Faculty of Agriculture, Niigata University, 2-8050

Ikarashi, Niigata 950-2181, Japan.

mnakano@agr.niigata-u.ac.jp

SOURCE: Scientia Horticulturae, (2002) Vol. 95, No. 1/2, pp.

123-132. 24 ref.

Publisher: Elsevier Science B.V. Amsterdam

ISSN: 0304-4238

DOI: 10.1016/S0304-4238(02)00033-X

PUB. COUNTRY: Netherlands Antilles

DOCUMENT TYPE: Journal LANGUAGE: English

ENTRY DATE: Entered STN: 20021206

Last Updated on STN: 20021206

AB Highly embryogenic callus cultures were established

from leaf explants in the Liliaceous ornamental plant, Agapanthus praecox subsp. orientalis, as the first step toward the development of an efficient transformation system. Embryogenic calluses were induced and then maintained by monthly subculturing onto a medium containing 1 mg picloram/litre. Upon transfer to a plant growth

regulator-free medium, the calluses produced numerous somatic embryos, most of which could develop into plantlets. Histological observations revealed that, following the transfer of the

observations revealed that, following the transfer of the embryogenic calluses to a plant growth regulator-free

medium, 2- to 6-cell proembryos, probably of unicellular origin, were produced, which passed through the globular and oval stages, and developed into club-shaped embryos with cotyledon, shoot apex and radicle. For establishing an efficient selection system in future transformation, the effects of selective agents (kanamycin, G418, hygromycin and bialaphos

[bilanafos]) and antibiotics for eliminating Agrobacterium (carbenicillin and cefotaxime) were examined on the growth and development of the embryogenic calluses. Callus growth

was completely inhibited by 50 mg hygromycin or 4 mg bialaphos/litre, and somatic embryo formation was completely inhibited by 50 mg hygromycin, 75 mg G418 or 3 mg bialaphos/litre. On the other hand, carbenicillin and cefotaxime promoted both growth and development of the embryogenic calluses.

L4 ANSWER 9 OF 11 CABA COPYRIGHT 2006 CABI on STN

ACCESSION NUMBER: 2003:30105 CABA DOCUMENT NUMBER: 20023198899

TITLE: Agrobacterium-mediated transformation in

Liliaceous ornamental plants

AUTHOR: Suzuki, S.; Nakano, M.

CORPORATE SOURCE: Faculty of Agriculture, Niigata University, 2-8050

Ikarashi, Niigata 950-2181, Japan.

mnakano@agr.niigata-u.ac.jp

SOURCE: JARQ, Japan Agricultural Research Quarterly, (2002)

Vol. 36, No. 3, pp. 119-127. 26 ref.

Publisher: Japan International Research Center for

Agricultural Sciences. Tsukuba

ISSN: 0021-3551

PUB. COUNTRY: Japan DOCUMENT TYPE: Journal English LANGUAGE:

Entered STN: 20030214 ENTRY DATE:

Last Updated on STN: 20030214

Studies on Agrobacterium-mediated transformation in 3 Liliaceous AB ornamental plants, Lilium formosanum, Agapanthus praecox ssp. orientalis and Muscari armeniacum, were described. Three different strains of A. tumefaciens were used, all of which harboured the binary vector carrying the nptII, hpt and qus-intron genes in the T-DNA region. For L. formosanum, no transgenic tissues nor plants were obtained after co-cultivation of organogenic calluses with A. tumefaciens, although transient expression of the gus gene could be detected in the calluses during co-cultivation. On the other hand, several hygromycin-resistant (Hygr) cell clusters were obtained for both A. praecox ssp. orientalis and M. armeniacum following the transfer of co-cultivated embryogenic calluses onto hygromycin (Hyg)-containing media. Hygr calluses developed into complete plants via somatic embryogenesis, and most of them were confirmed to be transgenic plants based on GUS histochemical assay and PCR analysis. Southern blot analysis revealed the integration of 1 to 5 copies of the transgene into the genome of the transgenic plants of both 2 species, but most of them had 1 or 2 copies. Agrobacterium -mediated transformation systems developed for A. praecox ssp. orientalis and M. armeniacum may be useful as a tool for their genetic improvement as well as molecular biology studies.

ANSWER 10 OF 11 CABA COPYRIGHT 2006 CABI on STN L4

ACCESSION NUMBER: 2001:104509 CABA

DOCUMENT NUMBER: 20013082803

Production of transgenic plants of the Liliaceous TITLE:

ornamental plant Agapanthus praecox ssp. orientalis

(Leighton) Leighton via Agrobacterium -mediated transformation of embryogenic

calli

Suzuki, S.; Supaibulwatana, K.; Mii, M.; Nakano, M.; AUTHOR:

Kanyaratt Supaibulwatana

Faculty of Agriculture, Niigata University, 2-8050 CORPORATE SOURCE:

Ikarashi, Niigata 950-2181, Japan.

Plant Science, (2001) Vol. 161, No. 1, pp. 89-97. 29 SOURCE:

ref.

Publisher: Elsevier Science Ltd. Oxford

ISSN: 0168-9452 United Kingdom

PUB. COUNTRY: DOCUMENT TYPE: Journal

English LANGUAGE:

Entered STN: 20011004 ENTRY DATE:

Last Updated on STN: 20011004

A system for producing transgenic plants was developed for the Liliaceous AB ornamental A. praecox ssp. orientalis via Agrobacterium-mediated genetic transformation. Leaf-derived embryogenic calluses were inoculated with A. tumefaciens strain EHA101/pIG121Hm or LBA4404/pTOK233, both of which harbored the binary vector carrying the neomycin phosphotransferase II (NPTII), hygromycin phosphotransferase (HPT) and intron-containing [beta]-glucuronidase (GUS-intron) genes in the T-DNA region. Following co-cultivation, the calluses were transferred to a medium containing 1 mg 1-1 picloram (PIC), 50 mg l-1 hygromycin and 500 mg l-1 cefotaxime, on which several hygromycin-resistant (Hygr) cell clusters were obtained 5-6 weeks after transfer. Agrobacterium strain, co-cultivation period and acetosyringone (AS) treatment during co-cultivation affected the number of Hygr callus lines produced: the best result was obtained when

embryogenic calluses were co-cultivated with

LBA4404/pTOK233 for 7 days in the presence of 20 mg l-1 AS. Hygr calluses were transferred to the same medium, but lacking PIC, for inducing somatic embryos. Somatic embryos thus obtained developed into complete plantlets following their transfer to a medium without PIC and antibiotics. All of them were verified to be stable transformants by GUS histochemical assay, PCR and Southern blot analyses.

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ANSWER 11 OF 11 CAPLUS

L4

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2000:790242 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         133:330528
                         Transformation of Allium sp. with
TITLE:
                         agrobacterium using embryogenic
                         callus cultures
INVENTOR(S):
                         Reynolds, John
PATENT ASSIGNEE(S):
                         Seminis Vegetable Seeds, Inc., USA
SOURCE:
                         PCT Int. Appl., 22 pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                         KIND
                                 DATE
                                             APPLICATION NO.
                                                                    DATE
                                 20001109
     WO 2000065903
                          A1
                                             WO 2000-US12463
                                                                     20000505
             AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU,
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             IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA,
             MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI,
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         RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
             DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
             CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     EP 1180927
                          A1
                                 20020227
                                          EP 2000-932149
                                                                    20000505
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                                 20051221
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO
     AU 780954
                          B2
                                 20050428
                                             AU 2000-49918
                                                                    20000505
PRIORITY APPLN. INFO.:
                                             US 1999-132617P
                                                                    19990505
                                             WO 2000-US12463
                                                                    20000505
AB
     The present invention relates to a method for transforming Allium
     species with a heterologous gene using Agrobacterium.
REFERENCE COUNT:
                               THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS
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=> d his
     (FILE 'HOME' ENTERED AT 15:29:27 ON 19 JAN 2006)
     FILE 'CAPLUS, CABA, AGRICOLA, BIOSIS' ENTERED AT 15:29:36 ON 19 JAN 2006
            221 S ALLI? AND AGROBACT?
L1
          13141 S EMBRYOGEN? AND CALL?
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             12 S L1 AND L2
             11 DUP REM L3 (1 DUPLICATE REMOVED)
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'FISTULOSUM) ' IS NOT A VALID FORMAT
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in at least one of the files. Refer to file specific help messages
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or the STNGUIDE file for information on formats available in

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individual files.
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 In a multifile environment, a format can only be used if it is valid
 in at least one of the files. Refer to file specific help messages
 or the STNGUIDE file for information on formats available in
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 L1
 AN
      2005:1268989 CAPLUS
      The efficacy of a novel insecticidal protein, Allium sativum
 TI
      leaf lectin (ASAL), against homopteran insects monitored in transgenic
      tobacco
      Dutta, Indrajit; Saha, Prasenjit; Majumder, Pralay; Sarkar, Anindya;
 ΑU
      Chakraborti, Dipankar; Banerjee, Santanu; Das, Sampa
      Plant Molecular and Cellular Genetics, Bose Institute, Kolkata, 700054,
 CS
      India
      Plant Biotechnology Journal (2005), 3(6), 601-611
 SO
      CODEN: PBJLAE; ISSN: 1467-7644
      Blackwell Publishing Ltd.
 PB
      Journal
 DT
      English
 LA
 RE.CNT 38
               THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD
               ALL CITATIONS AVAILABLE IN THE RE FORMAT
 => s l1 and (cepa or fistulosum)
            105 L1 AND (CEPA OR FISTULOSUM)
 L5
 => s 15 and 12
 L6
              4 L5 AND L2
- => dup rem 16
 PROCESSING COMPLETED FOR L6
 L7
               3 DUP REM L6 (1 DUPLICATE REMOVED)
 => d 1-3
      ANSWER 1 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 1
 L7
 AN
      2003:933684 CAPLUS
      140:175930
 DN
      Transgenic rose lines harboring an antimicrobial protein gene, Ace-AMP1,
 TI
      demonstrate enhanced resistance to powdery mildew (Sphaerotheca pannosa)
      Li, Xiangqian; Gasic, Ksenjia; Cammue, Bruno; Broekaert, Willem; Korban,
 ΑU
      Schuyler S.
      Department of Natural Resources and Environmental Sciences, University of
 CS
      Illinois, Urbana, IL, 618001, USA
 SO
      Planta (2003), 218(2), 226-232
      CODEN: PLANAB; ISSN: 0032-0935
 PB
      Springer-Verlag
 DT
      Journal
      English
 LA
 RE.CNT 34
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 L7
      ANSWER 2 OF 3 CAPLUS
                             COPYRIGHT 2006 ACS on STN
 AN
      2002:977962 CAPLUS
 DN
      138:36240
      Improved efficiency of regeneration of transgenic plants using
 TI
      meristematic or nodal tissue transformed with Agrobacterium
      Goldman, Stephen L.; Rudrabhatla, Sairam V.
 IN
      University of Toledo, USA
 PΑ
 SO
      PCT Int. Appl., 84 pp.
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CODEN: PIXXD2
DT
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     English
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FAN.CNT 1
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     PATENT NO.
                                                                    DATE
                        A2
     WO 2002102979
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                                20040729
     WO 2002102979
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             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,
             UA, UG, US, UZ, VN, YU, ZA, ZM, ZW
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     CA 2451863
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                                            EP 2002-742106
     EP 1455568
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     US 2004237133
                        A1
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                                            US 2003-480865
                                                                    20031212
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PRAI US 2001-298542P
                                20010615
     US 2002-356563P P
                                20020211
     WO 2002-US18966
                     W
                                20020614
     ANSWER 3 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN
L7
AN
     2000:790242
                CAPLUS
DN
     133:330528
     Transformation of Allium sp. with agrobacterium using
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     embryogenic callus cultures
IN
     Reynolds, John
PA
     Seminis Vegetable Seeds, Inc., USA
     PCT Int. Appl., 22 pp.
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     CODEN: PIXXD2
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     Patent
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     EP 1180927
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RE.CNT 3
              THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
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(FILE 'HOME' ENTERED AT 15:29:27 ON 19 JAN 2006)

FILE 'CAPLUS, CABA, AGRICOLA, BIOSIS' ENTERED AT 15:29:36 ON 19 JAN 2006 221 S ALLI? AND AGROBACT? L1L213141 S EMBRYOGEN? AND CALL? L312 S L1 AND L2 11 DUP REM L3 (1 DUPLICATE REMOVED) L4L5 105 S L1 AND (CEPA OR FISTULOSUM) L6 4 S L5 AND L2 3 DUP REM L6 (1 DUPLICATE REMOVED) L7 => d 17 1-3 ibib abs ANSWER 1 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 1 L7 2003:933684 CAPLUS ACCESSION NUMBER: 140:175930 DOCUMENT NUMBER: Transgenic rose lines harboring an antimicrobial TITLE: protein gene, Ace-AMP1, demonstrate enhanced resistance to powdery mildew (Sphaerotheca pannosa) AUTHOR (S): Li, Xiangqian; Gasic, Ksenjia; Cammue, Bruno; Broekaert, Willem; Korban, Schuyler S. Department of Natural Resources and Environmental CORPORATE SOURCE: Sciences, University of Illinois, Urbana, IL, 618001, USA Planta (2003), 218(2), 226-232 SOURCE: CODEN: PLANAB; ISSN: 0032-0935 Springer-Verlag PUBLISHER: DOCUMENT TYPE: Journal English LANGUAGE: An antimicrobial protein gene, Ace-AMP1, was introduced into Rosa hybrida AB cv. Carefree Beauty via Agrobacterium-mediated transformation. A total of 500 putative transgenic plants were obtained from 100 primary embryogenic calli co-cultivated with A. tumefaciens following selection on a regeneration medium containing 100 mg/l kanamycin. Polymerase chain reaction anal. of these putative transgenic lines, using primers for both Ace-AMP1 and neomycin phosphotransferase (npt II) genes, showed that 62% of these plants were pos. for both transgenes. lines were further confirmed for stable integration of Ace-AMP1 and npt II genes by Southern blotting. Transcription of the Ace-AMP1 transgene in various transgenic rose lines was determined using Northern blotting. Transgenic rose lines inoculated with conidial spores of Sphaerotheca pannosa (Wallr.: Fr.) Lev. var. rosae showed enhanced resistance to powdery mildew using both a detached-leaf assay and an in vivo greenhouse whole-plant assay. REFERENCE COUNT: 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 2 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN L7 2002:977962 CAPLUS ACCESSION NUMBER: 138:36240 DOCUMENT NUMBER: Improved efficiency of regeneration of transgenic TITLE: plants using meristematic or nodal tissue transformed with Agrobacterium Goldman, Stephen L.; Rudrabhatla, Sairam V. INVENTOR(S): PATENT ASSIGNEE(S): University of Toledo, USA PCT Int. Appl., 84 pp. SOURCE: CODEN: PIXXD2 DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

KIND DATE

APPLICATION NO.

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PATENT NO.

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WO 2002-US18966
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                                            WO 2002-US18966
     Methods of efficiently transforming monocotyledonous and dicotyledonous
AB
     plant tissue and regenerating plants with a very high yield of transgenic
     plants are described. The method uses Agrobacterium to
     transform root or apical meristem that is then cultured under conditions
     that generate somatic embryogenesis. The time required for the
     production of transgenic plants is significantly decreased, while the number of
     transgenic plants is significantly increased. These increases are not
     dependent upon the use of super-virulent Agrobacterium strains.
     The invention also relates to an improved technique for in vitro
     regeneration of mono- and di-cotyledonous plants in a suitable medium
     containing a novel growth regulator regime that promotes cell elongation in
     the production of numerous somatic embryos that are regenerable into fertile
     plants. Optimization expts. for the transformation of grasses and legumes
     using a \beta-glucuronidase reporter gene are described. Efficient
     genotype-independent regeneration of transgenic corn is demonstrated.
     ANSWER 3 OF 3
                    CAPLUS COPYRIGHT 2006 ACS on STN
L7
                         2000:790242 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         133:330528
                         Transformation of Allium sp. with
TITLE:
                         agrobacterium using embryogenic
                         callus cultures
                         Reynolds, John
INVENTOR(S):
PATENT ASSIGNEE(S):
                         Seminis Vegetable Seeds, Inc., USA
SOURCE:
                         PCT Int. Appl., 22 pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
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                         English
LANGUAGE:
FAMILY ACC. NUM. COUNT:
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PATENT INFORMATION:
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PATENT NO. KIND DATE APPLICATION NO. DATE

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EP 1180927 B1 20051221

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IE, SI, LT, LV, FI, RO

AU 780954 B2 20050428 AU 2000-49918 20000505 PRIORITY APPLN. INFO.: US 1999-132617P P 19990505

WO 2000-US12463 W 20000505

AB The present invention relates to a method for transforming Allium species with a heterologous gene using Agrobacterium.

REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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 => s onion and agrobact?
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      ANSWER 1 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN
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      2003:933684 CAPLUS
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      Transgenic rose lines harboring an antimicrobial protein gene, Ace-AMP1,
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      demonstrate enhanced resistance to powdery mildew (Sphaerotheca pannosa)
      Li, Xiangqian; Gasic, Ksenjia; Cammue, Bruno; Broekaert, Willem; Korban,
 AU
      Schuyler S.
      Department of Natural Resources and Environmental Sciences, University of
 CS
      Illinois, Urbana, IL, 618001, USA
      Planta (2003), 218(2), 226-232
 SO
      CODEN: PLANAB; ISSN: 0032-0935
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      Journal
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 RE.CNT 34
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      ANSWER 2 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN
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      Transformation of Allium sp. with agrobacterium using
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      embryogenic callus cultures
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ANSWER 3 OF 3 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved. (2006) on STN

AN 2004:10606 AGRICOLA

DN IND43618018

TI Transgenic rose lines harboring an antimicrobial protein gene, Ace-AMP1, demonstrate enhanced resistance to powdery mildew (Sphaerotheca pannosa).

AU Li, X.; Gasic, K.; Cammue, B.; Broekaert, W.; Korban, S.S.

AV DNAL (450 P693)

SO Planta, 2003 Dec. Vol. 218, no. 2 p. 226-232 ISSN: 0032-0935

NTE Includes references

DT Article

FS Non US

LA English